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What is claimed is:

- 1. A complex comprising:
- a folate receptor alpha (FRA)-expressing cancer cell from a subject; and

bound thereto, a conjugate comprising:

- an Fc portion of an IgA antibody, wherein said Fc portion comprises a CH2 domain and a CH3 domain but does not comprise a hinge region; and folate or a folate analog.
- 2. The complex of claim 1, wherein the conjugate further comprises a linker.
- 3. The complex of claim 2, wherein the linker is a peptide linker
- **4**. The complex of claim **2**, wherein the linker is a non-peptide linker.
- 5. The complex of claim 1, wherein the Fc portion is a variant Fc.
- **6**. The complex of claim **1**, wherein the Fc portion is a variant Fc and the conjugate comprises folate.
- 7. The complex of claim 1, wherein the Fc portion is a variant Fc and the conjugate comprises a folate analog.
- **8**. The complex of claim **1**, wherein the Fc fragment consists of a CH2/CH3 region from an IgA antibody.
- 9. The complex of claim 1, which is in vitro, or in vivo in the subject.
- 10. A method of inducing antibody-dependent cell-mediated cytotoxicity (ADCC), against folate receptor alpha (FRA)-expressing cancer cells, in neutrophils in a subject, the method comprising:
 - administering to the subject a therapeutically effective amount of a conjugate comprising:
 - an Fc portion of an IgA antibody, wherein said Fc portion comprises a CH2 domain and a CH3 domain but does not comprise a hinge region; and

folate or a folate analog,

- thereby triggering ADCC, against FRA-expressing cancer cells, in neutrophils in the subject.
- 11. The method of claim 10, wherein the cancer cells are breast cancer, ovarian cancer, or lung cancer cells.
- 12. The method of claim 10, wherein the cancer cells are triple negative breast cancer cells.
- 13. The method of claim 10, wherein the cancer cells that lack receptors for one or more of estrogen, progesterone and epithelial growth factor.
- 14. The method of claim 10, wherein administration is by intravenous administration.
- 15. The method of claim 10, wherein the conjugate is formulated for intravenous administration.